



# STIC Search Report

## Biotech-Chem Library

STIC Database Tracking Number: 151270

TO: Janet Epps-Ford  
Location: rem/2c05/2c18  
Art Unit: 1635

*April 29*, 2005

Case Serial Number: 10/086062

From: P. Sheppard  
Location: Remsen Building  
Phone: (571) 272-2529

sheppard@uspto.gov

### Search Notes

*04/29/05*

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STIC-Biotech/ChemLib

151270

me

From: Epps-Ford, Janet  
Sent: Wednesday, April 20, 2005 5:14 PM  
To: STIC-Biotech/ChemLib  
Subject: Sequence search.

Application 10/086,062

Please search SEQ ID NO: 4 in all pending and published nucleic acid databases.

Thanks,

Janet L. Epps-Ford, Ph.D.

Art Unit 1635

Mailbox: Remsen 2C18

Office: Remsen 2C05

Phone: 571-272-0757

Fax: 571-273-0757

\*\*\*\*\*

STAFF USE ONLY

Searcher: \_\_\_\_\_  
Searcher Phone: 2-\_\_\_\_\_  
Date Searcher Picked up: \_\_\_\_\_  
Date Completed: \_\_\_\_\_  
Searcher Prep/Rev. Time: \_\_\_\_\_  
Online Time: \_\_\_\_\_

\*\*\*\*\*

Type of Search

NA#: \_\_\_\_\_ AA#: \_\_\_\_\_  
Interference: \_\_\_\_\_ SPDI: \_\_\_\_\_  
S/L: \_\_\_\_\_ Oligomer: \_\_\_\_\_  
Encode/Transl: \_\_\_\_\_  
Structure#: \_\_\_\_\_ Text: \_\_\_\_\_  
Inventor: \_\_\_\_\_ Litigation: \_\_\_\_\_

\*\*\*\*\*

Vendors and cost where applicable

STN: \_\_\_\_\_  
DIALOG: \_\_\_\_\_  
QUESTEL/ORBIT: \_\_\_\_\_  
LEXIS/NEXIS: \_\_\_\_\_  
SEQUENCE SYSTEM: \_\_\_\_\_  
WWW/Internet: \_\_\_\_\_  
Other(Specify): \_\_\_\_\_

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GenCore version 5.1.6  
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 26, 2005, 06:25:00 ; Search time 1680 Seconds  
(without alignments)  
865.272 Million cell updates/sec

Title: US-10-086-062-4  
Perfect score: 30  
Sequence: 1 ctggaccctctgcactgcagagttccgct 30

Scoring table: IDENTITY\_NUC  
Gapop 10.0, Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues  
Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : GenEmbl:  
1: gb\_ba:\*  
2: gb\_hcg:\*  
3: gb\_in:\*  
4: gb\_om:\*  
5: gb\_ov:\*  
6: gb\_pat:\*  
7: gb\_ph:\*  
8: gb\_pl:\*  
9: gb\_pr:\*  
10: gb\_ro:\*  
11: gb\_sts:\*  
12: gb\_sy:\*  
13: gb\_un:\*  
14: gb\_vl:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	30	100.0	30	6	AX358109 Sequence
2	20.4	68.0	595	9	HUMY076F01
3	20.4	68.0	98348	9	AL136103 Human DNA
4	20.4	68.0	178965	9	AC012048 Homo sapi
5	20.2	67.3	23211	3	AC099767 Caenorhab
6	20	66.7	1503	1	AB038027 Vibrio sp
7	20	66.7	156165	2	AC021453 Homo sapi
8	20	66.7	157980	9	AC100852 Homo sapi
9	19.6	65.3	257595	2	AC123011 Rattus no
10	19.4	64.7	11406	4	AF107201 Equus cab
11	19.4	64.7	70387	4	AC087075 Caenorhab
12	19.4	64.7	110000	8	AC104791 Homo sapi
13	19.4	64.7	159969	2	AC104791 Homo sapi
14	19.4	64.7	189269	2	AP006440 Homo sapi
15	19.4	64.7	196472	2	AC011862 Homo sapi
16	19.2	64.0	1500	1	AF118021 Vibrio ca
17	19.2	64.0	1500	6	AR452360 Sequence
18	19.2	64.0	1500	6	AX201179 Sequence
19	19.2	63.3	93695	8	ATF2009

c	20	19	63.3	156806	2	AC021353	AC021353 Homo sapi
c	21	19	63.3	177652	10	AC140251	AC140251 Mus muscu
c	22	19	63.3	199749	8	ARCHIV68	AL161572 Arabidops
c	23	19	63.3	203120	10	AC124991	AC124991 Mus muscu
c	24	19	63.3	205475	2	AC130033	AC130033 Rattus no
c	25	19	63.3	213906	2	AC118634	AC118634 Mus muscu
c	26	19	63.3	223675	9	AC025031	AC025031 Homo sapi
c	27	19	63.3	233320	10	AC127314	AC127314 Mus muscu
c	28	19	63.3	249974	2	AC111249	AC111249 Rattus no
c	29	18.8	62.7	2481	6	AB027828	AB027828 Sequence
c	30	18.8	62.7	2481	10	RATR1B2	M2366 Rattus norv
c	31	18.8	62.7	5550	1	AB016787	AB016787 Pseudomon
c	32	18.8	62.7	23107	6	AX695653	AX695653 Sequence
c	33	18.8	62.7	35758	2	AC127539	AC127539 Homo sapi
c	34	18.8	62.7	67145	2	AC124281	AC124281 Homo sapi
c	35	18.8	62.7	69833	2	AC010777	AC010777 Homo sapi
c	36	18.8	62.7	77691	9	AC120118	AC120118 Homo sapi
c	37	18.8	62.7	95241	9	HS39819	AL023096 Human DNA
c	38	18.8	62.7	100521	9	AC119397	AC119397 Homo sapi
c	39	18.8	62.7	101721	9	AC067950	AC067950 Homo sapi
c	40	18.8	62.7	110000	8	CR382131.15	Continuation (16'o
c	41	18.8	62.7	113196	9	HS0697K14	AL121829 Human DNA
c	42	18.8	62.7	141605	2	AC013732	AC013732 Homo sapi
c	43	18.8	62.7	146606	9	AC068213	AC068213 Homo sapi
c	44	18.8	62.7	148332	9	AC079586	AC079586 Homo sapi
c	45	18.8	62.7	148611	8	AC091680	AC091680 Oryza sat

## ALIGNMENTS

RESULT 1	AX358109	Sequence 4 from Patent WO0194394.	30 bp	DNA	linear	PAT 13-FEB-2002
LOCUS	AX358109					
DEFINITION	AX358109					
ACCESSION	AX358109.1	GI:18674856				
VERSION						
KEYWORDS						
SOURCE						
ORGANISM						
REFERENCE						
AUTHORS						
TITLE						
JOURNAL						
FEATURES						
source						
ORIGIN						
Query Match						
Best Local Similarity						
Matches						
Conservative						
Mismatches						
Indels						
Gaps						
RESULT 2						
HUMY076F01/c						
LOCUS						
DEFINITION						
ACCESSION						
VERSION						
KEYWORDS						
SOURCE						
ORGANISM						
Human sapiens						
Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;						

REFERENCE	TITLE	JOURNAL	REFERENCE	TITLE	JOURNAL	COMMENT
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.	1 (bases 1 to 595)	Moessner, J., Tan, F., Marra, M., Kucaba, T., Yandell, M., Martin, J., Marsh, G., Bowles, L., Wylie, T., Bowers, Y., Steptoe, M., Theising, B., Gelsel, S., Allen, M., Underwood, K., Chappell, J., Persson, B., Gibbons, M., Harvey, N., Pape, D., Chamberlain, A., Morales, R., Schurk, R., Ritter, E., Kohn, S., Swaller, T., Behrmer, K., Hillier, L., Wilson, R. and Waterston, R.	Submitted (24-AUG-1998)	Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA	Submitted BY: Genome Sequencing Center Department of Genetics Washington University St. Louis MO 63108, USA http://genome.wustl.edu/gsc mailto:est@wustl.wustl.edu	Full Clone Sequencing of the Longest Available Member from Each Unpublished 2 (bases 1 to 595) Waterston, R.
NOTICE: This sequence represents the full insert of this cDNA. No attempt has been made to verify whether this corresponds to the full-length of the original mRNA from which it was derived. We have tried to obtain double-stranded, or double chemistry sequence across the entire clone, but potentially, there are areas in the sequence where this level of coverage was not achieved. Nevertheless, we are confident of the accuracy of this sequence as all regions of low quality, as defined by PHRAP (P. Green, in preparation), were visually inspected and edited accordingly. The consensus quality values for this sequence have been submitted separately.						
FEATURES	Location of this clone is unknown.					
source	1..595					
	/organism="Homo sapiens"					
	/mol_type="mRNA"					
	/db_xref="taxon:9606"					
	/clone="IMAGE:201721"					
	/clone_lib="Scars_fetal_liver_spleen_1NPLS"					
	9..338					
	/rnc_family="L2"					
repeat_region						
ORIGIN						
Query Match	68.0%; Score 20.4; DB 9; Length 595;					
Best Local Similarity	80.0%; Pred. No. 2.9e+02;					
Matches	24; Conservative 0; Mismatches 6; Indels 0; Gaps 0;					
OR	1 CTGACCCCTCTCGACTCGAGAGTTCGCT 30					
	566 CTGACCCCTCTCGCTCGAGACTTCTCT 537					
RESULT 3	AL136103	98348 bp	DNA	linear	PRI 04-JAN-2001	
LOCUS	Human DNA sequence from clone RPL-250B11 on chromosome 10 contains					
DEFINITION	STS and GSS, complete sequence.					
ACCESSION	AL136103					
VERSION	AL136103.24					
KEYWORDS	HTG.					
SOURCE	Homo sapiens (human)					
ORGANISM	Homo sapiens					
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;					
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.					
TITLE	1 (bases 1 to 98348)					
JOURNAL	Submitted (04-JAN-2001)					
	Sanger Centre, Hinxton, Cambridgeshire,					

COMMENT

CB810 1SA. UK. E-mail enquiries: humquerry@sanger.ac.uk Clone requests: clone.requests@sanger.ac.uk

On Jul 28, 2000 this sequence version replaced gi:9501151.

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em', EMBL; Sw', SWISSPROT; Tr', TREMBL; Wp', WORMPEP. Information on the WORMPEP database can be found at [http://www.sanger.ac.uk/Projects/C\\_elegans/wormpep](http://www.sanger.ac.uk/Projects/C_elegans/wormpep) This sequence is the entire insert of clone RP1-250B11 This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated repeat sequence elements. Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key. This sequence was generated from part of bacterial clone contigs of human chromosome 10, constructed by the Sanger Centre Chromosome 10 Mapping Group.

Further information can be found at <http://www.sanger.ac.uk/HGP/Ch10>

RP1-250B11 is from the library RPC1-1 constructed by the group of Pieter de Jong. For further details see <http://www.choi1.org/bacpac/home.htm>

VECTOR: pCYPAC2.

FEATURES

source

1..98348

/organism="Homo sapiens"

/mol\_type="genomic DNA"

/db\_xref="taxon:9606"

/chromosome="10"

/clone="RP1-250B11"

/clone\_1fb="RPC1-1"

1..34

/note="Single clone region. Reads derived from clone PCR. Assembly consistent with restriction digest."

repeat\_region

2458..2765

/note="AluX repeat: matches 1..308 of consensus"

repeat\_region

3087..3267

/note="MER39b repeat: matches 355..546 of consensus"

repeat\_region

3269..3606

/note="MER39 repeat: matches 13..380 of consensus"

repeat\_region

6106..6337

/note="L2 repeat: matches 2435..2705 of consensus"

repeat\_region

6364..6662

/note="AluSg repeat: matches 1..298 of consensus"

repeat\_region

complement(8333..8799)

/note="match: GSS: Em:AQ0403443"

misc\_feature

complement(8375..8758)

/note="match: GSS: Em:AQ264393"

misc\_feature

8802..9234

/note="match: GSS: Em:AQ150662"

misc\_feature

9268..9356

/note="match: STS: Em:G28019"

repeat\_region

11809..12092

/note="AluX repeat: matches 1..284 of consensus"

repeat\_region

12205..12722

/note="L1ME repeat: matches 5285..5811 of consensus"

repeat\_region

14113..14171

/note="L2 repeat: matches 2692..2750 of consensus"

repeat\_region

14548..14846

/note="AluY repeat: matches 1..299 of consensus"

repeat\_region

15883..16169

/note="AluSX repeat: matches 1..287 of consensus"

repeat\_region

17146..18557

/note="MER52A repeat: matches 1..1755 of consensus"

repeat\_region

18559..18713

/note="MIR repeat: matches 6..160 of consensus"

repeat\_region

18955..19381

/note="MER3 repeat: matches 590..1061 of consensus"

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repeat_region 19384. .19519
/note="FLAM C repeat: matches 3. .137 of consensus"
repeat_region 19713. .19828
/note="MER63 repeat: matches 463. .577 of consensus"
repeat_region 19826. .20117
/note="MER63 repeat: matches 2. .307 of consensus"
repeat_region 20355. .20391
/note="MIR repeat: matches 88. .156 of consensus"
repeat_region 20440. .20701
/note="MIR repeat: matches 2. .248 of consensus"
misc_feature 22321. .22631
/note="match: STS: Em:HSU10C4"
misc_feature 25725. .26188
/note="match: GSS: Em:AQ211002"
misc_feature 25738. .25978
/note="match: GSS: Em:AQ823705"
misc_feature 26548. .27067
/note="match: GSS: Em:AQ798605"
misc_feature 26564. .27250
/note="match: GSS: Em:AQ055423"
misc_feature 27549. .27913
/note="match: GSS: Em:B87793"
repeat_region 28724. .28952
/note="L2 repeat: matches 2167. .2416 of consensus"
misc_feature complement(29565. .30037)
/note="match: GSS: Em:AQ155973"
misc_feature 30073. .30670
/note="match: GSS: Em:AQ195052"
misc_feature complement(31752. .32121)
/note="match: GSS: Em:AQ099305"
misc_feature complement(32344. .32928)
/note="match: GSS: Em:AQ485533"
misc_feature 32930. .33368
/note="match: GSS: Em:AQ424293"
repeat_region 34554. .34583
/note="15 copies 2 mer ca 100% conserved"
repeat_region 35059. .35135
/note="L2 repeat: matches 2417. .2500 of consensus"
misc_feature complement(35681. .36191)
/note="match: GSS: Em:B55923"
misc_feature complement(35777. .36102)
/note="match: GSS: Em:AQ223744"
misc_feature complement(35861. .36186)
/note="match: GSS: Em:AQ803876"
repeat_region 36692. .36800
/note="MIR repeat: matches 13. .123 of consensus"
repeat_region 40461. .40765
/note="AluX repeat: matches 1. .293 of consensus"
repeat_region 41312. .41502
/note="MIR repeat: matches 8. .192 of consensus"
repeat_region 41785. .41840
/note="28 copies 2 mer tg 85% conserved"
repeat_region 43022. .43334
/note="AluX repeat: matches 3. .312 of consensus"
misc_feature complement(44477. .45066)
/note="match: GSS: Em:AQ540223"
misc_feature complement(44671. .45063)
/note="match: GSS: Em:AQ337658"
repeat_region 45077. .45157
/note="MER57-internal repeat: matches 7151. .7230 of consensus"
repeat_region 45158. .45304
/note="LIP repeat: matches 5050. .5197 of consensus"
repeat_region 45284. .45621
/note="LIP repeat: matches 5476. .5812 of consensus"
misc_feature complement(45941. .46452)
/note="match: GSS: Em:AQ832143"
repeat_region 48564. .48709
/note="L2 repeat: matches 2596. .2749 of consensus"
repeat_region 48720. .49082
/note="THE1B repeat: matches 1. .364 of consensus"
repeat_region 49176. .49629
/note="MLT1D repeat: matches 1. .502 of consensus"

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repeat_region 49784. .49912
/note="FLAM A repeat: matches 8. .132 of consensus"
repeat_region 49914. .50164
/note="MIR repeat: matches 2. .261 of consensus"
repeat_region 52850. .52918
/note="L2 repeat: matches 2635. .2706 of consensus"
misc_feature 53092. .53613
/note="match: GSS: Em:AQ800606"
misc_feature 53110. .53608
/note="match: GSS: Em:AQ172250"
misc_feature 53151. .53604
/note="match: GSS: Em:AQ170181"
misc_feature 53165. .53586
/note="match: GSS: Em:AQ545768"
misc_feature 53179. .53355
/note="match: GSS: Em:AQ898990"
misc_feature 53378. .53834
/note="match: GSS: Em:AQ898990"
repeat_region 53776. .53958
/note="MIR repeat: matches 41. .250 of consensus"
misc_feature complement(53900. .54343)
/note="match: GSS: Em:AQ400632"
repeat_region 53959. .54203
/note="AluSq repeat: matches 65. .308 of consensus"
repeat_region 54204. .54216
/note="MIR repeat: matches 250. .260 of consensus"
misc_feature 54361. .54767
/note="match: GSS: Em:AQ729256"
repeat_region 54372. .54565
/note="match: GSS: Em:AQ729256"
misc_feature 54440. .54730
/note="match: GSS: Em:AQ084956"
misc_feature complement(55008. .55535)
/note="match: GSS: Em:AQ556262"
misc_feature 55610. .55935
/note="match: GSS: Em:AQ594863"
repeat_region 56871. .57293
/note="LMD repeat: matches 883. .1319 of consensus"
repeat_region 57297. .57602
/note="AluDo repeat: matches 1. .305 of consensus"

Query Match 68.0%; Score 20.4; DB 9; Length 98348;
Best Local Similarity 80.0%; Pred. No. 2e+02;
Matches 24; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

Qy 1 CTGAGCCCTCTCGACTGAGAGTTCGCT 30
Db 62365 CTGAGCCCTCTCGCTGAGACTTCCT 62394

RESULT 4
AC012048 178965 bp DNA linear PRI 20-APR-2002
LOCUS Homo sapiens chromosome 10 clone RP11-43N22, complete sequence.
DEFINITION AC012048
ACCESSION AC012048.11 GI:19744964
VERSION 1
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 178965)
AUTHORS Smith,D.R.
TITLE Genome Therapeutics Corporation Sequencing Center: Human Genome
Sequence Data
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 178965)
AUTHORS Smith,D.R.
TITLE Direct Submission
JOURNAL Submitted (19-OCT-1999) Genome Therapeutics Corporation, 100 Beaver
Street, Waltham, MA 02453, USA
REFERENCE 3 (bases 1 to 178965)
AUTHORS Smith,D.R.

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JOURNAL TITLE Direct Submission
Submitted (16-AUG-2001) Genome Therapeutics Corporation, 100 Beaver Street, Waltham, MA 02453, USA
REFERENCE AUTHORS Smith,D.R.
TITLE Direct Submission
JOURNAL Submitted (27-MAR-2002) Genome Therapeutics Corporation, 100 Beaver Street, Waltham, MA 02453, USA
REFERENCE AUTHORS Smith,D.R.
TITLE Direct Submission
JOURNAL Submitted (28-MAR-2002) Genome Therapeutics Corporation, 100 Beaver Street, Waltham, MA 02453, USA
COMMENT COMMENT On Mar 27, 2002 this sequence version replaced gi:15193325. FEATURES location/Qualifiers source .locusname="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606" /chromosome="10" /clone="RP11-43N22" /clone_lib="RPCI-11" ORIGIN Query Match 68.0%; Score 20.4; DB 9; Length 178965; * Best Local Similarity 80.0%; Pred.No.1.9e+02; Matches 24; Conservative 0; Mismatches 6; Indels 0; Gaps 0; QY 1 CTGAGCCCTCTCGACTCGAGAGTTCCGCT 30 ||| ||||| ||||| ||||| ||||| ||||| | Dg 32913 CTGAGCCCCCTCTCGCCTGAGACCTTCTCT 32942 ||| ||||| ||||| ||||| ||||| ||||| | RESULT 5 AC099767 23211 bp DNA linear INV 20-NOV-2001 LOCUS Caenorhabditis briggsae cosmid G24F01, complete sequence. DEFINITION AC099767 AC099767 VERSION AC099767.1 GI:17017639 HGX. SOURCE Caenorhabditis briggsae ORGANISM Caenorhabditis briggsae Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabdilitida; Rhabditoidea; Rhabdilitidae; Pelodicerinae; Caenorhabditis. REFERENCE AUTHORS Washington University Genome Sequencing Center. TITLE The C. briggsae Genome Sequencing Project JOURNAL Unpublished REFERENCE AUTHORS 2 (bases 1 to 23211) WATERSTON,R. TITLE Direct Submission JOURNAL Unpublished REFERENCE AUTHORS 3 (bases 1 to 23211) WATERSTON,R. TITLE Direct Submission SUBMITTED (20-NOV-2001) Department of Genetics, Washington University, Genome Sequencing Center, 444 Forest Park Avenue, St. Louis, MO 63110, USA COMMENTS Submitted by: Genome Sequencing Center Department of Genetics, Washington University St.Louis , MO 63110, USA email: rtw@nematoec.wustl.edu NOTICE: This sequence may not be the entire insert of this clone. It may be shorter because we only sequence overlapping sections once, or longer because we provide a small overlap between neighbouring submissions.
```

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This sequence was finished as follows unless otherwise noted: all
regions were double stranded, sequenced with an alternate chemistry
or covered by high quality data (i.e., phred quality >= 30); an
attempt was made to resolve all sequencing problems, such as
compressions and repeats; all regions were covered by sequence from
more than one m3 subclone.

location/Qualifiers
  1..23211
    /organism="Caenothabditis briggsae"
    /mol_type="genomic DNA"
    /strain="Gujarat G16"
    /db_xref="taxon:6238"

ORIGIN
Query Match      67.3%; Score 20.2; DB 3; Length 23211;
Best Local Similarity 86.0%; Pred. No. 2.7e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Y 4 GACCCCTCTCGACTCGAGAGTTCCG 28
   |||||
Db 14462 GATTCCTCTCACTCGAGAGTTCCG 14438

RESULT 6
LOCUS AB038027/c 1503 bp DNA linear BCT 18-FEB-2000
DEFINITION Vibrio sp. SG128 gene for 16S rRNA, partial sequence.
ACCESSION AB038027
VERSION AB038027.1 GI:7007473
KEYWORDS 16S ribosomal RNA.
SOURCE Vibrio sp. SG128
ORGANISM Vibrio sp. SG128
Bacteria: Proteobacteria; Gammaproteobacteria; Vibrionales;
Vibrionaceae; Vibrio.
1 (bases 1 to 1503)
Urakawa,H.
16S rRNA gene of marine bacterium
Published Only in DataBase (2000)
2 (bases 1 to 1503)
Urakawa,H.
Direct Submission
Submitted (05-FEB-2000) Hidetoshi Urakawa, Northwestern University,
Department of Civil Engineering; Technological Institute 2145
Sheridan Road, Evanston, Illinois 60208-3109, USA
(E-mail:h-urakawa@nwu.edu, Tel:+1-847-467-5710,
Fax:+1-847-491-4011)

FEATURES
      source
      location/Qualifiers
          1..1503
            /organism="Vibrio sp. SG128"
            /mol_type="genomic DNA"
            /strain="SG128"
            /db_xref="taxon:115126"
            <1..>1503
            /product="16S ribosomal RNA"

ORIGIN
Query Match      66.7%; Score 20; DB 1; Length 1503;
Best Local Similarity 82.1%; Pred. No. 4.1e+02;
Matches 23; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

Y 1 CTGAGCCCTCTCGACTCGAGAGTTCCG 28
   |||||
Db 231 CTGGGCCCATCCGACGCGAGAGTTCCG 204

RESULT 7
LOCUS AC021453/c 156165 bp DNA linear HTG 01-APR-2000
DEFINITION Homo sapiens clone RP11-125C16, WORKING DRAFT SEQUENCE, 14
unordered pieces.
ACCESSION AC021453
VERSION AC021453.3 GI:7382318

```



KEYWORDS HTG; HTGS PHASE1; HTGS\_DRAFT.

SOURCE Homo sapiens (human)

ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 156165)

AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.

TITLE Homo sapiens, clone RP11-125C16

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 156165)

AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N., Anderson, S., Baldwin, J., Barna, N., Beckert, R., Beda, F., Boguslavsky, L., Bouhgalter, B., Brown, A., Burkett, G., Caetle, A., Chepel, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P., Dekrellano, K., Dewar, K., Domino, M., Doyle, M., Fensholt, J., Ferreira, P., Fitzhugh, N., Forrest, C., Gage, D., Galagan, J., Gardyna, S., Grant, G., Hagos, B., Heald, A., Horton, L., Howland, J. C., Johnson, R., Jones, C., Kann, L., Karatas, A., Klein, J., Landers, T., Lehoczy, J., Levine, R., Liu, C., Liu, G., Locke, K., Macdonald, P., Margulis, N., McEwan, P., McGuirk, A., McKernan, K., McHeeters, R., Meldrum, J., Menene, L., Morrow, J., Naylor, J., Notman, C. H., O'Connor, T., O'Donnell, P., Ollivar, T. M., Peterson, K., Piere, N., Pisani, C., Pollara, V., Raymond, C., Riley, R., Rothman, D., Roy, A., Santos, R., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J., Titrrell, A., Vassiliev, H., Viel, R., Vo, A., Wu, X., Wyman, D., Ye, W. J., Zimmer, A. and Zody, M.

TITLE Direct Submission

JOURNAL Submitted (16-JAN-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

COMMENT On Apr 1, 2000 this sequence version replaced gi:6721267. All repeats were identified using RepeatMasker:

Smit, A. P. A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu

Project Information

Center project name: L5169

Center clone name: 125 C.16

Summary Statistics

Sequencing vector: M13; M77815; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 145142 bases at least Q40

Consensus quality: 151009 bases at least Q30

Consensus quality: 153345 bases at least Q20

Insert size: 16000; agarose-fp

Insert size: 154865; sum-of-contigs

Quality coverage: 4.1 in Q20 bases; agarose-fp

Quality coverage: 4.2 in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of 14 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 1836: contig of 1836 bp in length

1837 1936: gap of 100 bp

1937 5796: contig of 3860 bp in length

5797 5896: gap of 100 bp

5897 9797: contig of 3901 bp in length

9798 9897: gap of 100 bp

9898 13913: contig of 4022 bp in length

13920 14019: gap of 100 bp

14020 17400: contig of 3381 bp in length

17401 17500: gap of 100 bp

17501 21253: contig of 3753 bp in length

FEATURES

source

1. 156165

/organism="Homo sapiens"

/mol\_type="genomic DNA"

/db\_xref="taxon:9606"

/clone="RP11-125C16"

/clone\_id="RP11-125C16"

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/note="assembly\_fragment"

1937. 5796

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5897. 9797

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9898. 13919

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17501. 21253

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clone end:SP6

vector end:right

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31197. 41465

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/note="assembly\_fragment"

75348. 89874

/note="assembly\_fragment"

89974. gap of 100 bp

116362. contig of 26388 bp in length

116363 116463

156165: contig of 3703 bp in length.

Location/Qualifiers

21353: gap of 100 bp

21354 31096: contig of 9743 bp in length

31097 31196: gap of 100 bp

31197 41465: contig of 10269 bp in length

41466 41565: gap of 100 bp

41566 50901: contig of 9336 bp in length

51001: gap of 100 bp

51002 62041: contig of 11040 bp in length

62042 62141: gap of 100 bp

62142 75247: contig of 13106 bp in length

75248 75347: gap of 100 bp

75348 89874: contig of 14527 bp in length

89875 89974: gap of 100 bp

89975 116362: contig of 26388 bp in length

116363 116463

156165: contig of 3703 bp in length.

ORIGIN

Query Match 66.7%; Score 20; DB 2; Length 156165;

Best Local Similarity 82.1%; Pred. No. 2.9e+02;

Matches 23; Conservative 0; Mismatches 5; Indels 0;

2 TGGACCCCTCTCGACTCGAGAGTCCGC 29

11285 TGGACCTCTCTCTATTCAGAGTCTGC 11258

RESULT 8

LOCUS AC100852 157980 bp DNA linear PRI 29-AUG-2002

DEFINITION Homo sapiens chromosome 17, clone RP11-125C16, complete sequence.

ACCESSION AC100852

VERSION AC100852.2 GI:22539166

KEYWORDS HTG.

SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
AUTHORS Birren, B., Nussbaum, C. and Lander, E.  
TITLE 1 (bases 1 to 157980)  
JOURNAL Homo sapiens chromosome 17, clone RP11-125C16  
REFERENCE Unpublished  
AUTHORS 2 (bases 1 to 157980)  
Birren, B., Linton, L., Nussbaum, C., Lander, E., Ali, A., Allen, N.,  
Anderson, S., Barna, N., Bastien, V., Boguslavsky, L., Boukhgalter, B.,  
Brown, A., Camarata, J., Campolano, A., Chang, J., Chazaro, B.,  
Chapel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A.,  
Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S.,  
Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Garg, S.,  
Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-pierre, N.,  
Hagos, B., Heatford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R.,  
Jones, C., Kamat, A., Karatas, A., Kells, C., Lacroque, K.,  
Lamazares, R., Landers, T., Lehocsky, J., Levine, R., Liu, G.,  
Maclean, C., Macdonald, P., Major, J., Margus, N., Matthews, C.,  
McCarthy, M., McEwan, P., McKernan, K., McPheters, R., Meidrim, J.,  
Menues, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C.,  
Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D.,  
Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V.,  
Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P.,  
Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupack, R.,  
Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,  
Strauss, N., Subramaniam, A., Talamas, J., Tesfaye, S., Theodore, J.,  
Topham, K., Travers, M., Travis, N., Trigglio, J., Vassiliev, H.,  
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G.,  
Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.  
DIRECT SUBMISSION  
TITLE Submitted (22-NOV-2001) Whitehead Institute/MIT Center for Genome  
JOURNAL Research, 320 Charles Street, Cambridge, MA 02141, USA  
REFERENCE 3 (bases 1 to 157980)  
AUTHORS Birren, B., Nussbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,  
Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhgalter, B.,  
Camarata, J., Chang, J., Chazaro, B., Chappel, Y., Collymore, A.,  
Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S.,  
Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J.,  
Gardyna, S., Gord, S., Graham, L., Grand-pierre, N., Hagos, B.,  
Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A.,  
Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K.,  
Liu, G., Maclean, C., Macdonald, P., Major, J., Matthews, C.,  
McCarthy, M., Meidrim, J., Menues, L., Mihova, T., Mlenga, V.,  
Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H.,  
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,  
Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P.,  
Roman, J., Roy, A., Schauer, S., Schupack, R., Seaman, S., Severy, P.,  
Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J.,  
Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H.,  
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J.,  
Zembek, L., Zimmer, A. and Zody, M.  
DIRECT SUBMISSION  
TITLE Submitted (29-AUG-2002) Whitehead Institute/MIT Center for Genome  
JOURNAL Research, 320 Charles Street, Cambridge, MA 02141, USA  
COMMENT On Aug 29, 2002 this sequence version replaced g1:17048222.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1997)  
http://ftp.genome.washington.edu/RM/RepeatMasker.html  
----- Genome Center  
Center: Whitehead Institute/ MIT Center for Genome Research  
Center code: WITR  
Web site: http://www-seg.wi.mit.edu  
Contact: sequence\_submissions@genome.wi.mit.edu  
----- Project Information  
Center Project name: L21587  
Center clone name: 125\_C\_16  
----- Location/Qualifiers  
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Query Match      66.7%; Score 20; DB 9; Length 157980;
Best Local Similarity 82.1%; Pred. No. 2.9e+02;
Matches 23; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY      2 TGGACCCCTCGACTCGAGAGTCCGC 29
Db      130676 TGGACCTCTCTATCAAGAGTCTGC 130649

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RESULT 9
AC123011      257595 bp      DNA      linear      HTG 12-OCT-2002
LOCUS      Rattus norvegicus clone CH230-92124, WORKING DRAFT SEQUENCE, 4
DEFINITION      Rattus norvegicus (Norway rat)
ACCESSION      AC123011
VERSION      AC123011.3 GI:23665280
KEYWORDS      HTG: HTGS PHASE1; HTGS DRAFT; HTGS FULLTOP.
SOURCE      Rattus norvegicus
ORGANISM      Rattus norvegicus
            Eumetazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
            Rattus.
REFERENCE      1 (bases 1 to 257595)

```

## AUTHORS

Muzny D, Marie, Metzker M, Lee, Abramson S, Adams C, Alder J, Allen C, Allen H, Albrooks S, Amin A, Angiano D, Anyalebechi V, Aoyagi A, Ayodeji M, Baca E, Baden H, Baldwin D, Bandaranaike D, Barber M, Barnstead M, Bernhied F, Biewald K, Blair J, Blankenburg K, Blyth P, Brown M, Bryant N, Buhay C, Burch P, Barrell K, Calderon E, Cardenas V, Carter K, Cavazos I, Cesar H, Center A, Chacko J, Chavez D, Chen G, Chen R, Chen Y, Chen Z, Chu J, Cleveland C, Cockrell R, Cox C, Coyle M, Cree A, D Souza L, Davila M, L, Davis S, Davy-Carroll L, De Anda C, Dederich D, Delgado O, Denon S, Deramo C, Ding Y, Dinh H, Divya K, Draper H, Dugan-Rocha S, Dunn A, Durbin K, Duval B, Evans K, Egan A, Escotto M, Eugene C, Evans C, Falls T, Fan G, Fernandez S, Finley M, Flagg N, Forbes L, Foster M, Foster P, Fraser C, Gabisi A, Ganta R, Garcia A, Garner T, Garza M, Gebregeorgis E, Geer K, Gill R, Grady M, Guerra W, Guevara M, Gunaratne P, Haaland W, Hamill C, Hamilton C, Hamilton K, Harvey Y, Havlak P, Hawes A, Henderson N, Hernandez J, Hernandez R, Hines S, Hladun S, Hodgson A, Hogues M, Hollins B, Howells S, Hulys S, Hume J, Idledit D, Jackson A, Jackson L, Jacob L, Jiang H, Johnson B, Johnson R, Jolivet A, Karpachy S, Kelly S, Kelly S, Khan Z, King L, Kovar C, Kowis C, Kraft C, Lebow H, Levan J, Lewis L, Li Z, Liu J, Liu J, Liu W, Liu Y, London P, Longacre S, Lopez J, Lorenshewa L, Louised H, Lozano R, Lu X, Ma J, Maheshwari M, Mahindartne M, Mahmood M, Malloy K, Mangum A, Mangum B, Nagu P, Martin K, Martin R, Martinez E, Mawhiney S, McLeod M, McNeill T, Meenen E, Milosavljevic A, Miner G, Minja E, Montemayor J, Moore S, Morgan M, Morris K, Morris S, Munidasa M, Murphy M, Nair L, Nankervis C, Neal D, Newton N, Nguyen N, Norris S, Nwankwem O, Okunnu G, Olajunsgoon A, Pal S, Parks K, Pasternak S, Paul H, Perez A, Perez L, Plankoch C, Plapper F, Polidexter A, Popovic D, Primus E, Pu L, L, Puzo M, Quiroz J, Rachlin B, Reeves K, Regier M, A, Reigh R, Reilly B, Reilly M, Ren Y, Reuter M, Richards S, Riggs F, Rivers C, Rodley T, Rojia A, Rose M, Rose R, Ruiz S, J, Sanders W, Saverly G, Scherer S, Scott G, Shatman S, Shen H, Shetty J, Shvartsbeyn A, Sisson I, Siller C, D, Snajs D, Sneed A, Sodergren E, Song X, Z, Sorelle R, Sosa J, Steimle M, Strong R, Sutton A, Svatok A, Taber Z, Taylor C, Taylor T, Thomas N, Thomas S, Tingey A, Trejos Z, Usaml K, Valas R, Vera V, Villasana D, Waldron L, Walker B, Wang J, Wang Q, Wang S, Warren J, Warren R, Wei X, White F, Williams G, Willson R, Wleczky R, Wooden H, Worley K, Wright D, Wright R, Wu J, Yakub S, Yen J, Yoon L, Yoon V, Yu F, Zhang J, Zhou J, Zhou X, Zhao S, Dunn D, von Niederhausen A, Weiss R, Smith D, R, Holt R, A, Smith H, O, Weinstock G, and Gibbs R, A.

## TITLE

JOURNAL  
REFERENCE  
2 (bases 1 to 257595)

## AUTHORS

JOURNAL  
TITLE  
REFERENCE  
Rat Genome Sequencing Consortium.

## JOURNAL

## COMMENT

Submitted (12-OCT-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA  
On Oct 10, 2002 this sequence version replaced gi:21909149.  
The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole

genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

## ----- Genome Center

Center: Baylor College of Medicine  
Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: hgsc-help@bcm.tmc.edu

## ----- Project Information

Center project name: GYMD  
Center clone name: CH230-92124

## ----- Summary Statistics

Assembly program: Phrap; version 0.990329  
Consensus quality: 236342 bases at least Q40  
Consensus quality: 239870 bases at least Q30  
Consensus quality: 242005 bases at least Q20  
Estimated insert size: 243903; sum-of-coverage estimation  
Quality coverage: 7x in Q20 bases; sum-of-coverage estimation

\* NOTE: Estimated insert size may differ from sequence length  
\* (see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 4 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1 250677: contig of 250677 bp in length  
\* 250678 250777: gap of unknown length  
\* 250778 252307: contig of 1530 bp in length  
\* 252308 252407: gap of unknown length  
\* 252408 253577: contig of 1170 bp in length  
\* 253578 253678: gap of unknown length  
\* 253678 257595: contig of 3918 bp in length.

## FEATURES

## source

1. 257595  
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/mol\_type="genomic DNA"  
/db\_xref="taxon:10116"  
/clone="CH230-92124"  
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clone end:Sp6  
site:EcoRI  
end\_sequence:BH295371"  
misc\_feature  
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clone end:Sp6  
site:EcoRI  
end\_sequence:BH295371"

## ORIGIN

## Query Match

Best Local Similarity 84.6%; Pred. No. 4.2e+02;  
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CTGGACCCCTCTGAGCTCGAGAGTTC 26  
Db 151391 CTGGACCTCTTGAACCTTAGAGTTC 151416

RESULT 10  
AF107201/c 11406 bp DNA linear MAM 13-DEC-1998  
LOCUS AF107201  
DEFINITION Equus caballus beta-lactoglobulin II (BLG) gene, complete cds.  
ACCESSION AF107201  
VERSION AF107201.1 GI:4008111  
KEYWORDS  
SOURCE Equus caballus (horse)  
ORGANISM Equus caballus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

## REFERENCE

1 (bases 1 to 11406)  
Masel, A.M., Brandon, R.B. and Bell, T.K.  
Nucleotide Sequence of the Equine Beta-Lactoglobulin Gene  
JOURNAL  
TITLE  
AUTHORS  
TITLE  
JOURNAL

2 (bases 1 to 11406)  
Masel, A.M., Brandon, R.B. and Bell, T.K.  
Direct Submission  
Submitted (17-NOV-1998) Australian Equine Blood Typing Research  
Laboratory, University of Queensland, St. Lucia, Brisbane,  
Queensland 4072, Australia

## FEATURES

## source

1. 11406  
/organism="Equus caballus"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:9796"  
/gene="BLG"  
/gene="BLG"  
join(4472..4567,4956..5095,5767..5840,7367..7480,  
8302..8406,8663..8679)  
/product="beta-lactoglobulin II"  
/gene="BLG"  
join(4472..4567,4956..5095,5767..5840,7367..7480,  
8302..8406,8663..8679)  
/gene="BLG"  
/product="beta-lactoglobulin II"  
/codon\_start=1  
/protein\_id="AAC95385.1"  
/db\_xref="GI:4008112"

## CDS

translation="MKCLILALGSLMCGNQTDTPTQMODDLDGVAGRMHVSVA  
SDISLSDSSVPLRYVVEELRPTPEGNLIIIRGNHACVEENVAQTEDEAVFTV  
NYGKERKISVLDITVAHVMFPCVGPPLPSAHEGMQYLAFTQKDEVEWKEKRSALQ  
PLPGRVQIVQDPSCGGERCGF"

## ORIGIN

## Query Match

Best Local Similarity 79.3%; Pred. No. 6.5e+02;  
Matches 23; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 CTGGACCCCTCTGAGCTCGAGAGTTCGC 29  
Db 6919 CGGGGCCCTCTTGAAGTGAAGTTACAC 6891

## RESULT 11

## AC087075

AC087075 70387 bp DNA linear INV 05-DEC-2000  
Caenorhabditis briggsae cosmid CB023K10, complete sequence.

## DEFINITION

## ACCESSION

## VERSION

## KEYWORDS

## SOURCE

## ORGANISM

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## TITLE

## AUTHORS

## JOURNAL

## COMMENT

Submitted by: Department of Genetics, Washington  
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA  
Genome Sequencing Center  
Department of Genetics, Washington University,  
St. Louis, MO 63110, USA  
e-mail: jspliet@watson.wustl.edu

NOTICE: This sequence may not be the entire insert of this clone.  
It may be shorter because we only sequence overlapping sections

```

/chromosome="4"
/map="4"
/clone="RP11-181K12"
/clone_lib="RP11-11"
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2333..2362
/rpt_family="AT_rich"
2471..2690
/rpt_family="MIR"
3143..3169
/rpt_family="(A)n"
3296..3316
/rpt_family="AT_rich"
3304..3462
/rpt_family="Alu"
3589..3609
/rpt_family="AT_rich"
3984..4033
/rpt_family="ERV1"
4019..4111
/rpt_family="(TA)n"
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5195..5483
/rpt_family="Alu"
5374..5402
/rpt_family="(GGATG)n"
5897..5927
/rpt_family="(TTTTG)n"
5906..6217
/rpt_family="Alu"
6577..6760
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(NID:g7937452)"
7129..7131
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(NID:g7937452)"
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15896..15941
/rpt_family="(TG)n"
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16918..16939
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18098..18220

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21764..22061
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21780..22172
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23268..23312
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23637..24312
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24463..24756
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24798..25023
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25377..25406
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Query Match 64.7%; Score 19.4; DB 9; Length 159969;  
 Best Local Similarity 79.3%; Pred. No. 5.3e+02;  
 Matches 23; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

Qy 2 TGAGCCCTCGACTCGAGAGTTCCGCT 30  
 Db 120906 TGAGCTACTCTCGCTCCAGACTTCTCT 120878

RESULT 14  
 LOCUS AP006440/c 189269 bp DNA linear HTG 21-MAY-2003  
 DEFINITION Homo sapiens chromosome 11 clone RP11-368120 map 11g, WORKING DRAFT  
 ACCESSION AP006440  
 VERSION AP006440.1 GI:30962586  
 KEYWORDS HTG; HTGS PHASE1; HTGS\_DRAFT.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1  
 Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P.,  
 Fujiyama, A., Yada, T., Tokoki, Y., Watanabe, H. and Sakaki, Y.  
 Homo sapiens genomic DNA of 11g  
 Published Only in Database (2003)  
 2 (bases 1 to 189269)  
 Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P.,  
 Fujiyama, A., Yada, T., Tokoki, Y., Watanabe, H. and Sakaki, Y.  
 Direct Submission  
 Submitted (19-MAY-2003) Masahira Hattori, The Institute of Physical  
 and Chemical Research (RIKEN), Genomic Sciences Center (GSC),  
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
 (E-mail: hattori@gsc.riken.go.jp, URL: http://hgp.gsc.riken.go.jp/  
 Tel: 81-45-503-9111, Fax: 81-45-503-9170)

once, or longer because we provide a small overlap between neighboring submissions.

FEATURES  
source  
1. 70387  
/organism="Caenorhabditis briggsae"  
/mol\_type="genomic DNA"  
/strain="Gujarat G16"  
/db\_xref="taxon:6238"  
/clone="CB023K10"  
23139. 23209  
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complement(36569. 36640)  
/product="rRNA-Glu"  
/note="codon recognized: GAG"  
37437. 37508  
/product="rRNA-Gln"  
/note="codon recognized: CAA"

trna

trna

trna

ORIGIN

Query Match 64.7% Score 19.4; DB 3; Length 70387;  
Best Local Similarity 79.3% Pred. No. 5.7e+02;  
Matches 23; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 2 TGAACCCCTCTGACTGAGAGTTCCGCT 30  
Db 6252 TGGACACGCTCTGATTCGAAAGTTGAGCT 6280

RESULT 12

AB016816\_6/c  
WPCOMMENT

Sequence split into 9 fragments LOCUS AB016816 Accession AB016816

Fragment Name	Begin	End
AB016816_1	1	110000
AB016816_2	20001	310000
AB016816_3	300001	410000
AB016816_4	400001	510000
AB016816_5	500001	610000
AB016816_6	600001	710000
AB016816_7	700001	810000
AB016816_8	800001	907057

Continuation (7 of 9) of AB016816 from base 600001 (AB016816 Eremothecium gossypii ATCC

Query Match 64.7% Score 19.4; DB 8; Length 110000;  
Best Local Similarity 79.3% Pred. No. 5.5e+02;  
Matches 23; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 CTGACCCCTCTGACTGAGAGTTCCGC 29  
Db 20976 CTGACGCTGCTCGATCGAGAGTTCCGC 20948

RESULT 13

AC104791/c

LOCUS AC104791 Homo sapiens BAC clone RP11-181K12 from 4, complete sequence.  
DEFINITION AC104791 AC032008  
ACCESSION AC104791.3 GI:18482313  
VERSION  
KEYWORDS  
HTG.  
SOURCE  
ORGANISM Homo sapiens (human)  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
PUBMED  
REFERENCE  
AUTHORS

1 (bases 1 to 159969)  
Toward a complete human genome sequence  
Genome Res. 8 (11), 1097-1108 (1998)  
9847074  
2 (bases 1 to 159969)  
Isak, A., Meyer, R. and Creason, K.

TITLE  
JOURNAL  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
Submitted (21-DEC-2001) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA  
4 (bases 1 to 159969)  
Waterston, R.H.  
Direct Submission  
Submitted (03-FEB-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA  
5 (bases 1 to 159969)  
Waterston, R.H.  
Direct Submission  
Submitted (21-FEB-2002) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA  
On Feb 3, 2002 this sequence version replaced gi:18042374.

COMMENT

Center: Washington University Genome Sequencing Center  
Center code: WUGSC  
Web site: <http://genome.wustl.edu/gsc>  
Contact: [sapiens@watson.wustl.edu](mailto:sapiens@watson.wustl.edu)  
----- Summary Statistics  
Center project name: H\_NH0181K12  
Drafting Center: W18R

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:  
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:  
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:  
The RP11-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Moon, P.Y., Zhao, B., Frengen, E., Tareno, M., Catanesse, J.J., and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at <http://www.chori.org>  
VECTOR: pBAC3.6

NEIGHBORING SEQUENCE INFORMATION:  
The clone sequenced to the left is RP11-292M9, the clone sequenced to the right is RP11-203B7. Actual start of this clone is at base position 1 of RP11-181K12; actual end is at base position 159969 of RP11-181K12.

A transposon has been identified in the vector of this clone.

The sequence of AC032008 has been incorporated into AC104791.

FEATURES  
source

1. 159969  
/organism="Homo sapiens"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:9606"



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11826 12707: contig of 882 bp in length  
12708 12807: gap of 100 bp  
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13698 13797: gap of 100 bp  
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19535 19634: gap of 100 bp  
19635 20510: contig of 876 bp in length  
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20611 21517: contig of 907 bp in length  
21518 21617: gap of 100 bp  
21617 22499: contig of 882 bp in length  
22500 22599: gap of 100 bp  
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23481 23580: gap of 100 bp  
23581 24444: contig of 864 bp in length  
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32426 32525: gap of 100 bp  
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36371 36470: gap of 100 bp  
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37459 38332: contig of 874 bp in length  
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54025 54124: gap of 100 bp  
54125 54957: contig of 833 bp in length  
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60896 61774: contig of 879 bp in length  
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62840 63807: contig of 968 bp in length  
63808 63907: gap of 100 bp  
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64784 64883: gap of 100 bp  
64884 65767: contig of 884 bp in length  
65768 65867: gap of 100 bp  
65869 66765: contig of 898 bp in length  
66766 66865: gap of 100 bp  
66866 67730: contig of 865 bp in length  
67731 67830: gap of 100 bp  
67831 68721: contig of 891 bp in length  
68722 68821: gap of 100 bp  
68822 69711: contig of 890 bp in length  
69712 69811: gap of 100 bp  
69812 70699: contig of 888 bp in length  
70699 70799: gap of 100 bp  
70799 71679: contig of 880 bp in length  
71679 70800

Query Match 64.7%; Score 19.4; DB 2; Length 196472;  
Best Local Similarity 79.3%; Pred. No.5,je+02;  
Matches 23; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 CTGAGCCCTCTGCACTCGAGAGTTCCGC 29  
Db 6946 CTGAGCCCTCTGCACTCGAGAGTTCCGC 6918

Search completed: April 26, 2005, 11:39:38  
Job time : 1686 secs



GenCore version 5.1.6  
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 26, 2005, 06:19:21 ; Search time 427 Seconds  
(without alignments)  
415.907 Million cell updates/sec

Title: US-10-086-062-4  
Perfect score: 30  
Sequence: 1 ctgaccctctcgcacgcagagctcgcgc 30

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2359870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : N\_Geneseq\_16Dec04:  
1: geneseqn1980s:\*  
2: geneseqn1990s:\*  
3: geneseqn2000s:\*  
4: geneseqn2001s:\*  
5: geneseqn2001bs:\*  
6: geneseqn2002as:\*  
7: geneseqn2002bs:\*  
8: geneseqn2003as:\*  
9: geneseqn2003bs:\*  
10: geneseqn2003cs:\*  
11: geneseqn2003ds:\*  
12: geneseqn2004as:\*  
13: geneseqn2004bs:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	30	100.0	30	AAD24069	Aad24069 Maize eng
2	19.2	64.0	1500	AAS11024	Aas11024 Vibrio ch
3	19	63.3	121124	AD097107	Adg97107 Mouse can
4	18.8	62.7	2481	ABT41766	ABT41766 Toxicity
5	18.8	62.7	23107	ADA02762	Ada02762 Human RUN
6	18.8	62.7	23107	ADB72500	Adb72500 Human Run
7	18.8	62.7	23107	ADC85242	Adc85242 Human Run
8	18.8	62.7	23107	ADM74357	Adm74357 Human car
9	18.6	62.0	3210	AAZ27624	Aaz27624 Plasmid S
10	18.6	62.0	3459	AAZ27623	Aaz27623 Plasmid S
11	18.4	61.3	319608	AAH51601	Aah51601 Human chr
12	18.4	61.3	319608	AAH51601	Aah51601 Human chr
13	18.2	60.7	340449	AAH52198	Aah52198 Human sch
14	18	60.0	42325	ADM74382	Adm74382 Mycobacte
15	18	60.0	110000	ADM74382	Adm74382 Mycobacte
16	17.8	59.3	165	ADD49400	Add49400 Human lun
17	17.8	59.3	254	ADG00322	Adg00322 Nicotiana
18	17.8	59.3	369	ABD11207	Abd11207 Pseudomon
19	17.8	59.3	426	ADD49385	Add49385 Human lun
20	17.8	59.3	449	ADD49343	Add49343 Human lun

## ALIGNMENTS

21	17.8	59.3	449	10	ADD49294	Add49294 Human lun
22	17.8	59.3	986	6	ABQ46720	Abq46720 Oligonuc1
23	17.8	59.3	986	6	ABQ46721	Abq46721 Oligonuc1
24	17.8	59.3	1553	12	ADL12873	Adl12873 Human ste
25	17.8	59.3	1603	4	AAK77077	Aak77077 Human lun
26	17.8	59.3	1731	11	ACN91401	Acn91401 Breast ca
27	17.8	59.3	2780	13	ADR25834	Adr25834 Breast ca
28	17.8	59.3	152141	8	ACA64961	Ac64961 Human BCR
29	17.8	59.3	153995	13	ABD33534	Abd33534 Murine ca
30	17.6	58.7	246	12	AD052469	Ad052469 Human met
31	17.6	58.7	8119	3	AAZ35392	Aaz35392 Maize ete
32	17.6	58.7	49243	4	ABL03188	AbL03188 Drosoph11
33	17.4	58.0	65	6	ABN51822	Abn51822 Mouse sp1
34	17.4	58.0	300	2	AAZ13036	Aaz13036 Human gen
35	17.4	58.0	300	2	AAZ98464	Aaz98464 Human can
36	17.4	58.0	411	4	AAI85652	Aai85652 Human pol
37	17.4	58.0	460	9	ACH41661	Ach41661 Human foe
38	17.4	58.0	594	12	ACH74968	Ach74968 Human gen
39	17.4	58.0	699	2	AAZ15926	Aaz15926 Human gen
40	17.4	58.0	729	12	ADO63441	Ado63441 Transcript
41	17.4	58.0	778	6	ABK30430	Abk30430 Human G-p
42	17.4	58.0	936	13	ADR92877	Adr92877 Novel S.
43	17.4	58.0	949	13	ADS05077	Ads05077 Bacterial
44	17.4	58.0	1210	4	AA41038	Aa41038 CDNA enco
45	17.4	58.0	1210	4	AA34819	Aa34819 CDNA enco

## RESULT 1

AAD24069 standard; DNA; 30 BP.

AAD24069;

09-APR-2002 (first entry)

Maize engineered Ubi-1 promoter heat shock element #3.

Gene expression; maize; ubiquitin promoter; Ubi-1; HSE;

heat shock element; agronomic gene; ds.

Zea mays.

Key Location/Qualifiers

misc\_feature 1..15

FT /note= "5' heat shock element"

FT /tag= b

FT /note= "3' heat shock element"

WT 0200194394-A2.

13-DEC-2001.

08-JUN-2001; 2001WO-US018689.

09-JUN-2000; 2000US-00590558.

(PROD-) PRODIGENE INC.

Jilka JM, Hood EE, Howard JA;

WPI; 2002-122117/16.

New promoter sequences for causing expression of a structural gene especially agronomic gene or open reading frame in a plant cell, comprises engineered versions of the maize ubiquitin promoter.

Claim 6; Page 54; 68pp; English.

CC The invention relates to a promoter sequence capable of directing  
CC expression of a nucleotide sequence in a plant cell, comprising maize  
CC ubiquitin (ubi-1) promoter sequence with a modification so that it does  
CC not include two overlapping heat shock elements (HSE) or it directs  
CC expression to increase the endosperm/embryo expression ratio of the  
CC protein when compared to the ratio from a wild-type ubiquitin promoter.  
CC The modified ubi-1 promoter comprises a deletion of 3', 5' or both HSEs,  
CC two non-overlapping/adjacent HSEs, replacement of HSEs with a trimer of a  
CC seed specific element from the promoter of pea lectin gene Psi, or  
CC insertion of a transcription factor binding site in the HSE region. An  
CC expression construct comprising modified ubi-1 promoter is useful for  
CC causing expression of a structural gene (agronomic genes) or open reading  
CC frame in a plant cell. The modified ubi-1 promoter increases expression  
CC levels beyond those observed with native ubiquitin promoter. The present  
CC sequence is maize engineered ubi-1 promoter with heat shock elements  
CC adjacent placed. Note: The present sequence is also shown in claim 26,  
CC page 56 of the specification. However, this sequence has an additional  
CC nucleotide at the 3' end

SO Sequence 30 BP; 4 A; 12 C; 7 G; 7 T; 0 U; 0 Other;  
Query Match 100.0%; Score 30; DB 6; Length 30;  
Best Local Similarity 100.0%; Pred. No. 0.0012;  
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CTGAGCCCTCTCGACTCGAGATTCCGCT 30  
DB 1 CTGAGCCCTCTCGACTCGAGATTCCGCT 30

## RESULT 2

AAS1024/c  
AAS1024 standard; DNA; 1500 BP.

AC AAS1024;

DT 11-SEP-2003 (revised)

DT 24-OCT-2001 (first entry)

DE Vibrio cholera 16S ribosomal RNA gene.

XX Antisense; bacterial 16S ribosomal RNA; rRNA; bacterial infection; human;  
KM food grain supplement; livestock; poultry; therapeutic; ds.

OS Vibrio cholerae.

PN WO200142457-A2.

PD 14-JUN-2001.

PF 29-NOV-2000; 2000WO-US042391.

PR 29-NOV-1999; 99US-0168150P.

PA (AVIB-) AVI BIOPHARMA INC.

PI Iversen PL;

DR WPI; 2001-457295/49.

PT Antibacterial compound, useful for treating bacterial infections and as  
PT livestock and poultry food supplement, comprises antisense  
PT oligonucleotides complementary to bacterial 16S and 23S rRNA.

PS Disclosure; Page; 62pp; English.

XX AAS1021-AAS1024 represent the coding sequences of bacterial 16S  
CC ribosomal RNA (rRNA) genes. The sequences were used to design anti-  
CC bacterial compounds comprising substantially unchanged antisense  
CC nucleic acid sequence at least 10 nucleotides in length which is  
CC complementary to a bacterial 16S or 23S rRNA nucleic acid sequence. The  
CC antisense oligomers are used for treating a bacterial infection in a

CC human or a mammalian animal produced by Escherichia coli, Salmonella  
CC typhimurium, Pseudomonas aeruginosa, Vibrio cholera, Neisseria  
CC gonorrhoea, Helicobacter pylori, Bartonella henselae, Haemophilus  
CC influenzae, Shigella dysenteriae, Staphylococcus aureus, Mycobacterium  
CC tuberculosis, Streptococcus pneumoniae, Treponema pallidum and Chlamydia  
CC trachomatis. The antibacterial compound may be used as a food grain  
CC supplement in livestock and poultry food composition. Note: The present  
CC sequence is not shown in the specification but has been accessed from  
CC Genbank using the appropriate accession number given in the  
CC specification. (Updated on 11-SEP-2003 to standardise OS field)

SO Sequence 1500 BP; 376 A; 326 C; 482 G; 312 T; 0 U; 4 Other;  
Query Match 64.0%; Score 19.2; DB 5; Length 1500;  
Best Local Similarity 75.0%; Pred. No. 1.1e+02;  
Matches 21; Conservative 2; Mismatches 5; Indels 0; Gaps 0;  
QY 1 CTGAGCCCTCTCGACTCGAGATTCCG 28  
DB 212 CTGGGCCCATCCGACGCGARARGTCG 185

## RESULT 3

ADQ97107/c  
ADQ97107 standard; DNA; 121124 BP.

AC ADQ97107;

DT 07-OCT-2004 (first entry)

DE Mouse cancer associated sequence MD08-002, SEQ ID 83.

XX Cytostatic; Gene Therapy; cancer; leukemia; lymphoma; Mouse; ds.

OS Mus musculus.

PN WO2004060304-A2.

PD 22-JUL-2004.

PF 22-DEC-2003; 2003WO-US041389.

PR 27-DEC-2002; 2002US-00330773.

PA (SAGR-) SAGRES DISCOVERY INC.

PI Morris DW, Malandro MS;

DR WPI; 2004-543781/52.

PT New isolated cancer associated nucleic acids comprising at least 10  
PT contiguous nucleotides, useful for diagnosing, preventing and/or treating  
PT cancers such as leukemia and lymphoma.

PS Claim 1; SEQ ID NO 83; 199pp; English.

CC The present invention relates to cancer associated sequences (ADQ97025-  
CC ADQ98004). The sequences are useful for the diagnosis, prevention and/or  
CC treatment of cancer, such as leukemia and lymphoma. Note: The sequence  
CC data for this patent did not form part of the printed specification, but  
CC was obtained in electronic format directly from WIPO at  
CC ftp.wipo.int/pub/published\_pct\_sequences.

SO Sequence 121124 BP; 32972 A; 25314 C; 26641 G; 35451 T; 0 U; 746 Other;

Query Match 63.3%; Score 19; DB 12; Length 121124;  
Best Local Similarity 81.5%; Pred. No. 1.9e+02;  
Matches 22; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 4 GACCCCTCTCGACTCGAGATTCCGCT 30  
DB 88764 GCCCCTCTCGATTACAGAGTTCCGCT 88738

RESULT 4
ABT41766/C
ID ABT41766 standard; DNA; 2481 BP.
XX AC
XX DT
DT 26-JUN-2003 (first entry)
DE Toxicity modelling related rat gene SEQ ID No 1468.
XX DE
XX KW
KW Toxic effect; gene expression profile; renal toxicity; toxicity marker; database; drug screening; toxicity assay; rat; ds.
XX OS
OS Rattus norvegicus.
PN WO200295000-A2.
XX PD
PD 28-NOV-2002.
XX PE
PE 22-MAY-2002; 2002MO-US016173.
XX PF
PF 22-MAY-2001; 2001US-0292335P.
PR 13-JUN-2001; 2001US-0297523P.
PR 19-JUN-2001; 2001US-0298925P.
PR 10-JUL-2001; 2001US-0303807P.
PR 10-JUL-2001; 2001US-0303808P.
PR 10-JUL-2001; 2001US-0303810P.
PR 28-AUG-2001; 2001US-0315047P.
PR 27-SEP-2001; 2001US-0324928P.
PR 22-OCT-2001; 2001US-0330462P.
PR 01-NOV-2001; 2001US-0330867P.
PR 21-NOV-2001; 2001US-0331805P.
PR 06-DEC-2001; 2001US-0336144P.
PR 19-DEC-2001; 2001US-0340873P.
PR 21-FEB-2002; 2002US-0357842P.
PR 21-FEB-2002; 2002US-0357843P.
PR 21-FEB-2002; 2002US-0357844P.
PR 15-MAR-2002; 2002US-0364134P.
PR 08-APR-2002; 2002US-0370144P.
PR 08-APR-2002; 2002US-0370206P.
PR 08-APR-2002; 2002US-0370247P.
PR 17-APR-2002; 2002US-0372794P.
PR 21-APR-2002; 2002US-0371679P.
XX PA
PA (GENE-) GENE LOGIC INC.
P1 Mendrick D, Porter M, Johnson K, Higgs B, Castle A, Elashoff M;
XX WPI; 2003-148464/14.
DR XX
PT Predicting at least one toxic effect of a compound, useful for toxicity modelling, comprises preparing a gene expression profile of a tissue or cell sample exposed to the compound, and comparing the gene expression profile to a database.
XX PS
PS Example 4; Page; 446pp; English.
XX CC
CC The invention relates to a novel method of predicting at least one toxic effect of a compound. The method comprises a gene expression profile of a tissue or cell sample exposed to the compound, and comparing the gene expression profile to a database comprising at least part of the data or information given in the specification. The methods are useful for predicting at least one toxic effect of a compound, predicting the progression of a toxic effect of a compound, predicting the renal toxicity of a compound, or identifying toxicity markers in tissues or cells exposed to known renal toxin. The genes are useful as toxicity markers in drug screening and toxicity assays, in monitoring disease or physiological states, or disease progression. This polynucleotide represents a rat DNA sequence relating to the toxic effect database described in the specification. NOTE: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from the World Intellectual Property

CC Organization  
SQ Sequence 2481 BP; 599 A; 653 C; 555 G; 674 T; 0 U; 0 Other;  
  
Query Match 62.7%; Score 18.8; DB 10; Length 2481;  
Best Local Similarity 76.7%; Pred. No. 1.7e+02;  
Matches 23; Conservative 0; Mismatches 7; Indels 0; Gaps 0;  
  
QY 1 CTGACCCCTCTCGACTCGAGATTCCGCT 30  
DB 1077 CTGTCCCCCTCTCGTCTAGGGATTCAGCT 1048  
  
RESULT 5  
ADA02762 ID ADA02762 standard; DNA; 23107 BP.  
ADA02762 AC  
XX ADA02762;  
XX  
DT 06-NOV-2003 (first entry)  
DE Human RUNX3 carcinoma associated gene, SEQ ID NO:1280.  
XX  
KV Human; carcinoma associated; oncogene; carcinoma; cancer; breast;  
KW prostate; lymphoma; leukaemia; cytostatic; gene therapy; drug screening;  
gene; ds.  
XX  
OS Homo. sapiens.  
XX  
PN WO2003057146-A2.  
PD 17-JUN-2003.  
PF 26-DEC-2002; 2002WO-USO41414.  
PR 26-DEC-2001; 2001US-00035832.  
PA (SAGR-) SAGRES DISCOVERY.  
PI Morris DW;  
DR WPI; 2003-587068/55.  
PT New recombinant nucleic acid encoding carcinoma associated protein,  
useful for preparing compositions for treating carcinomas.  
PS Claim 1; SEQ ID NO 1280; 245dp; English.  
XX  
XX The invention relates to recombinant carcinoma associated (CA) nucleic  
acid sequences from mouse and human (ADA01482-ADA03094), and to  
recombinant carcinoma associated proteins (CAR) encoded by them. The  
invention also encompasses expression vectors and host cells comprising a  
CA nucleic acid, a polypeptide (especially an antibody) that specifically  
binds to the protein, and a biochip comprising CA nucleic acid or  
fragments thereof. The sequences of the invention were identified using  
oncogenic retroviruses, which insert into the genome of the host organism  
at random. Many of these do not carry transduced host oncogenes or  
pathogenic trans-acting viral genes, meaning that cancer incidence is a  
direct consequence of the effects of proviral integration into host  
protooncogenes. The CA nucleic acid sequences can be used to diagnose  
carcinoma (especially breast cancer, prostate cancer, lymphoma or  
leukemia) or a propensity to carcinoma by determination of the sequence  
of a CA gene, or by determination of CA gene expression in particular  
tissues. CA nucleic acid, proteins and antibodies are also useful as  
therapeutic agents and in screening and evaluating drug candidates. The  
present sequence represents a specifically claimed human CA nucleic acid  
sequence of the invention. Note: The complete sequence data for this  
c patent did not form part of the printed specification, but was obtained  
in electronic format directly from WIPO at  
ftp.wipo.int/pub/published\_pct\_sequences.

Sequence 23107 BP; 5308 A; 6264 C; 6229 G; 5306 T; 0 U; 0 Other;

Query Match 62.7%; Score 18.8; DB 9; Length 23107;  
 Best Local Similarity 76.7%; Pred. No. 2.1e+02;  
 Matches 23; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 1 CTGGACCCCTCTCGACTCGAGATTCCGCT 30  
 14873 CTGGAGAGCCCTGACTCCAGATTCCGCT 14902

## RESULT 6

ADB72500  
 ID ADB72500 standard; DNA; 23107 BP.

AC ADB72500;

DT 04-DEC-2003 (first entry)

DE Human Runx3 gene.

human; ds; cytostatic; gene therapy; vaccine; carcinoma; lymphomas;  
 cancer; neoplasm; adenocarcinoma; sarcoma; gene.

OS Homo sapiens.

PN W02003008583-A2.

PD 30-JAN-2003.

PF 26-DEC-2001; 2001WO-US051291.

PR 02-MAR-2001; 2001US-00798586.

PR 23-OCT-2001; 2001US-00004113.

PR 08-NOV-2001; 2001US-00052482.

PR 30-NOV-2001; 2001US-00997722.

PR 20-DEC-2001; 2001US-00034650.

PA (SAGR-) SAGRES DISCOVERY.

PI Morris DW, Engelhard EK;

PT WPI; 2003-239337/23.

PS Claim 1; SEQ ID NO 328; 2304pp; English.

CC The invention relates to a novel recombinant nucleic acid comprising a

CC nucleotide sequence selected from any of the 660 sequences fully defined

CC in the specification. A polynucleotide of the invention has cytostatic

CC activity, and may have a use in gene therapy, or in a vaccine. The

CC recombinant nucleic acids and polypeptides are useful for treating

CC carcinomas, e.g. lymphomas, cancers, neoplasm, adenocarcinoma, and

CC sarcomas. The present sequence represents a human gene of the invention.

Sequence 23107 BP; 5308 A; 6264 C; 6229 G; 5306 T; 0 U; 0 Other;

QY 1 CTGGACCCCTCTCGACTCGAGATTCCGCT 30  
 14873 CTGGAGAGCCCTGACTCCAGATTCCGCT 14902

## RESULT 7

ADCB5242  
 ID ADCB5242 standard; DNA; 23107 BP.

AC ADCB5242;

DT 01-JAN-2004 (first entry)

XX Human Runx3 genomic sequence.

DE Cytostatic; gene therapy; vaccine; cancer; carcinoma-associated gene; CA;

KM secreted; transmembrane; intracellular; ds.

OS Homo sapiens.

PN W02003045230-A2.

PD 05-JUN-2003.

PF 02-DEC-2002; 2002WO-US038582.

PR 30-NOV-2001; 2001US-00997722.

PA (SAGR-) SAGRES DISCOVERY.

PI Morris DW, Engelhard EK;

PT WPI; 2003-513603/48.

PS New recombinant nucleic acid comprising a nucleotide sequence of any of

the carcinoma-associated (CA) genes, useful for screening for drug

candidates for diagnosing or treating carcinomas.

Claim 1; SEQ ID NO 28; 983pp; English.

CC The invention relates to a recombinant nucleic acid comprising a

CC nucleotide sequence selected from any of the fully defined carcinoma-

CC associated (CA) genes from the 50 tables given in the specification. The

CC CA proteins are secreted, transmembrane or intracellular proteins. The

CC recombinant nucleic acids are useful for screening for drug candidates

CC for diagnosing or treating carcinomas. Sequences given in ADCB5215-

CC ADCB5514 represent CA genes of the invention.

Sequence 23107 BP; 5308 A; 6264 C; 6229 G; 5306 T; 0 U; 0 Other;

QY 1 CTGGACCCCTCTCGACTCGAGATTCCGCT 30  
 14873 CTGGAGAGCCCTGACTCCAGATTCCGCT 14902

ADCB5242  
 ID ADCB5242 standard; DNA; 23107 BP.

AC ADCB5242;

DT 01-JUN-2004 (first entry)

DE Human carcinoma associated (CA) nucleic acid #13.

XX Human; carcinoma associated nucleic acid; CA nucleic acid; gene; ds;

KM carcinoma associated protein; CAP; carcinoma; leukemia; lymphoma;

OS Cytostatic.

PN US2004072154-A1.

PD 15-APR-2004.

PF 30-NOV-2001; 2001US-00997722.

PR 22-DEC-2000; 2000US-00747377.

PR 02-MAR-2001; 2001US-00798586.

PA (MORR/) MORRIS D W.

PA (ENGELHARD E. K.  
 XX Morris DW, Engelhard EK;  
 PI WPI, 2004-328562/30.  
 DR  
 XX New carcinoma associated gene or protein, useful for preparing a  
 PT composition for diagnosing or treating carcinoma e.g., leukemia or  
 PT lymphoma.  
 XX  
 PS Claim 1; SEQ ID NO 28; 29pp; English.  
 XX  
 CC The invention relates to new recombinant nucleic acids. The invention  
 CC also relates to a host cell comprising a recombinant nucleic acid or  
 CC expression vector, an expression vector comprising a recombinant nucleic  
 CC acid, a recombinant protein, a method of screening for drug candidates, a  
 CC method of screening for a bioactive agent capable of binding to a  
 CC carcinoma associated protein (CAP) encoded by a nucleotide sequence, a  
 CC method of screening for a bioactive agent capable of modulating the  
 CC activity of a CAP, a method of evaluating the effect of a candidate  
 CC carcinoma drug, a method of diagnosing carcinoma, a method for inhibiting  
 CC the activity of a CAP, a method of treating carcinomas, a method of  
 CC neutralising the effect of a CAP and a method of diagnosing carcinoma or  
 CC propensity to carcinoma. A method of evaluating the effect of a candidate  
 CC carcinoma drug comprises administering the drug to a patient, removing a  
 CC cell sample from the patient and determining alterations in the  
 CC expression or activation of a gene comprising the nucleotide sequence. A  
 CC method of diagnosing carcinoma comprises determining the expression of  
 CC one or more genes comprising the nucleic acid sequence in a first tissue  
 CC type of a first individual and comparing the expression of the gene from  
 CC a second normal tissue type from the first individual or a second  
 CC unaffected individual, where a difference in the expression indicates  
 CC that the first individual has carcinoma. A method of inhibiting the  
 CC activity of a CAP comprises binding an inhibitor to the CAP. Treating  
 CC carcinomas comprises administering to a patient an inhibitor of CAP.  
 CC Neutralising the effect of a CAP comprises contacting an agent specific  
 CC for the CAP. The polypeptide specifically binds to the protein encoded by  
 CC the nucleic acid. It comprises an antibody that specifically binds to the  
 CC protein encoded by the nucleic acid. The nucleic acids are useful for  
 CC preparing a composition for diagnosing or treating carcinoma e.g.,  
 CC leukemia or lymphoma. This sequence represents a human carcinoma  
 CC associated (CA) nucleic acid of the invention. Note: The sequence data  
 CC for this patent did not form part of the printed specification but was  
 CC obtained in electronic format directly from USPTO at  
 CC seqdata.uspto.gov/sequence.html.  
 CC  
 XX  
 CC  
 SQ Sequence 23107 BP; 5308 A; 6264 C; 6229 G; 5306 T; 0 U; 0 Other;  
 Query Match 62.7%; Score 18.8; DB 12; Length 23107;  
 Best Local Similarity 76.7%; Pred. No. 2.1e+02;  
 Matches 23; Conservative 0; Mismatches 7; Indels 0; Gaps 0;  
 QY 1 CTGAGCCCTCTCGACTCGAGAGTTCCGCT 30  
 DB 14873 CTGGAGGCCCTCGACTCCAGATTCGCT 14902  
 RESULT 9  
 AA227624/c  
 ID AA227624 standard; DNA; 3210 BP.  
 XX  
 AC AA227624;  
 XX  
 DT 20-DEC-1999 (first entry)  
 XX  
 DE Plasmid SP1F-1.  
 XX  
 KW Extracellular compartment modification; floral cell; self-compatibility;  
 KW pollen-pistil interaction; self-incompatibility; insect growth control;  
 KW plasmid SPF-1; GP1S363 gene; cysteine protease inhibitor gene; ss.  
 XX  
 OS Synthetic.  
 XX

PN WO9949063-A1.  
 XX  
 PD 30-SEP-1999.  
 XX  
 PF 19-MAR-1999; 99WO-CA000237.  
 XX  
 PR 20-MAR-1998; 98US-0078728P.  
 XX  
 PA (MIAC ) CANADA MIN AGRIC & AGRI-FOOD CANADA.  
 XX  
 PI Robert LS, Gledzie S;  
 XX  
 DR WPI, 1999-591104/50.  
 XX  
 PT Protein expression in floral cells for peptide display, mediating plant  
 PT sterility, and modifying pollen-pistil interactions.  
 XX  
 PS Example 6; Page 103-104; 113pp; English.  
 XX  
 CC This sequence represents the plasmid SP1F-1, containing a fusion of the  
 CC Brassica napus GP1S363 gene to the Onchocerca protease inhibitor gene.  
 CC The invention relates to a method for modifying the extracellular  
 CC compartment of a floral cell of a plant, that comprises expressing a  
 CC construct comprising a gene of interest encoding a protein, fusion  
 CC protein or peptide, or a fragment of them, which is capable of modifying  
 CC the composition of the extracellular compartment of the floral cell and  
 CC altering either the function, use or development of the floral cell or  
 CC modifying the interaction of the floral cell with other cells, within an  
 CC anther or pistil cell. The method is used to modify pollen-pistil  
 CC interaction or function, which mediates, produces or prevents self-  
 CC compatibility, self-incompatibility, out- or in-crossing or combinations  
 CC of these. The method is also used for localizing proteins on the surface  
 CC of pollen for the purpose of peptide display. The protein localized on  
 CC the surface of the pollen may be an antibody or antigen or is a protein  
 CC that is effective in controlling insect growth, behaviour, feeding,  
 CC development or reproduction  
 CC  
 XX  
 SQ Sequence 3210 BP; 1050 A; 626 C; 595 G; 939 T; 0 U; 0 Other;  
 Query Match 62.0%; Score 18.6; DB 2; Length 3210;  
 Best Local Similarity 84.0%; Pred. No. 2.2e+02;  
 Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 1 CTGAGCCCTCTCGACTCGAGAGTT 25  
 DB 1232 CTGAGCCCTTCAATGATGATT 1208  
 RESULT 10  
 AA227623/c  
 ID AA227623 standard; DNA; 3459 BP.  
 XX  
 AC AA227623;  
 XX  
 DT 20-DEC-1999 (first entry)  
 XX  
 DE Plasmid SPF-1.  
 XX  
 KW Extracellular compartment modification; floral cell; self-compatibility;  
 KW pollen-pistil interaction; self-incompatibility; insect growth control;  
 KW plasmid SPF-1; GP1S363 gene; cysteine protease gene; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO9949063-A1.  
 XX  
 PD 30-SEP-1999.  
 XX  
 PF 19-MAR-1999; 99WO-CA000237.  
 XX  
 PR 20-MAR-1998; 98US-0078728P.  
 XX  
 PA (MIAC ) CANADA MIN AGRIC & AGRI-FOOD CANADA.  
 XX  
 PI Robert LS, Gledzie S;  
 XX  
 DR WPI, 1999-591104/50.  
 XX  
 PT Protein expression in floral cells for peptide display, mediating plant  
 PT sterility, and modifying pollen-pistil interactions.  
 XX  
 PS Example 6; Page 103-104; 113pp; English.  
 XX  
 CC This sequence represents the plasmid SP1F-1, containing a fusion of the  
 CC Brassica napus GP1S363 gene to the Onchocerca protease inhibitor gene.  
 CC The invention relates to a method for modifying the extracellular  
 CC compartment of a floral cell of a plant, that comprises expressing a  
 CC construct comprising a gene of interest encoding a protein, fusion  
 CC protein or peptide, or a fragment of them, which is capable of modifying  
 CC the composition of the extracellular compartment of the floral cell and  
 CC altering either the function, use or development of the floral cell or  
 CC modifying the interaction of the floral cell with other cells, within an  
 CC anther or pistil cell. The method is used to modify pollen-pistil  
 CC interaction or function, which mediates, produces or prevents self-  
 CC compatibility, self-incompatibility, out- or in-crossing or combinations  
 CC of these. The method is also used for localizing proteins on the surface  
 CC of pollen for the purpose of peptide display. The protein localized on  
 CC the surface of the pollen may be an antibody or antigen or is a protein  
 CC that is effective in controlling insect growth, behaviour, feeding,  
 CC development or reproduction  
 CC  
 XX  
 SQ Sequence 3210 BP; 1050 A; 626 C; 595 G; 939 T; 0 U; 0 Other;  
 Query Match 62.0%; Score 18.6; DB 2; Length 3210;  
 Best Local Similarity 84.0%; Pred. No. 2.2e+02;  
 Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 1 CTGAGCCCTCTCGACTCGAGAGTT 25  
 DB 1232 CTGAGCCCTTCAATGATGATT 1208  
 RESULT 10  
 AA227623/c  
 ID AA227623 standard; DNA; 3459 BP.  
 XX  
 AC AA227623;  
 XX  
 DT 20-DEC-1999 (first entry)  
 XX  
 DE Plasmid SPF-1.  
 XX  
 KW Extracellular compartment modification; floral cell; self-compatibility;  
 KW pollen-pistil interaction; self-incompatibility; insect growth control;  
 KW plasmid SPF-1; GP1S363 gene; cysteine protease gene; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO9949063-A1.  
 XX  
 PD 30-SEP-1999.  
 XX  
 PF 19-MAR-1999; 99WO-CA000237.  
 XX  
 PR 20-MAR-1998; 98US-0078728P.  
 XX  
 PA (MIAC ) CANADA MIN AGRIC & AGRI-FOOD CANADA.  
 XX

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XX Robert LS, Gledlie S;
PI WPI; 1999-591104/50.
XX Protein expression in floral cells for peptide display, mediating plant
XX sterility, and modifying pollen-pistil interactions.
XX
PS Example 5; Page 100-102; 113pp; English.
XX
XX This sequence represents the plasmid SPF-1, containing a fusion of the
XX Brassica napus GPIS363 gene to the Sitophilus cysteine protease gene. The
XX invention relates to a method for modifying the extracellular compartment
XX of a floral cell of a plant, that comprises expressing a construct
XX comprising a gene of interest encoding a protein, fusion protein or
XX peptide, or a fragment of them, which is capable of modifying the
XX composition of the extracellular compartment of the floral cell and
XX altering either the function, use or development of the floral cell or
XX modifying the interaction of the floral cell with other cells, within an
XX anther or pistil cell. The method is used to modify pollen-pistil
XX interaction or function, which mediates, produces or prevents self-
XX compatibility, self-incompatibility, out- or in-crossing or combinations
XX of these. The method is also used for localizing proteins on the surface
XX of pollen for the purpose of peptide display. The protein localized on
XX the surface of the pollen may be an antibody or antigen or is a protein
XX that is effective in controlling insect growth, behaviour, feeding,
XX development or reproduction
XX
SQ Sequence 3459 BP; 1082 A; 721 C; 678 G; 978 T; 0 U; 0 Other;
XX
XX Query Match 62.0%; Score 18.6; DB 2; Length 3459;
XX Best Local Similarity 84.0%; Pred. No. 2.2e+02;
XX Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
XX
QY 1 CTGAGCCCTCTCGACTCGAGACTT 25
DB 1232 CTGAGCCCTCTCGAGACTT 1208
XX
RESULT 11
AAH51601
XX AAH51601 standard; DNA; 319608 BP.
XX
AC AAH51601;
XX
XX 29-AUG-2001 (first entry)
XX
DE Human chromosome 13q31-q33 genomic nucleotide sequence.
XX
XX sbg1; g3465; sbg2; g35017; g35018; chromosome 13q31-q33; haplotype;
XX biallelic marker; polymorphism; schizophrenia; bipolar disorder; ds.
XX
OS Homo sapiens.
XX
PN WO200058510-A2.
XX
PD 05-OCT-2000.
XX
XX 30-MAR-2000; 2000MO-IB000435.
XX
XX 30-MAR-1999; 99US-0126903P.
XX 30-APR-1999; 99US-0131971P.
XX 30-APR-1999; 99US-0132065P.
XX 14-JUL-1999; 99US-0143928P.
XX 27-JUL-1999; 99US-0145915P.
XX 29-JUL-1999; 99US-0146452P.
XX 29-JUL-1999; 99US-0146453P.
XX 28-OCT-1999; 99US-0162288P.
XX
XX (GEST ) GENSET.
XX
XX Cohen D, Blumenfeld W, Chumakov I, Bougueleret L, Bihain B,
XX Bastoux L;
PI

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XX WPI; 2000-619082/59.
XX
XX Polynucleotides comprising sequences from sbg1 and g35018 biallelic
XX markers are used for genotyping and detecting schizophrenia or bipolar
XX disorder and predisposition to these disorders.
XX
PS Claim 1; Page 409-493; 737pp; English.
XX
XX AAH51601 represents a human genomic nucleotide sequence comprising sbg1,
XX g3465, sbg2, g35017 and g35018 nucleic acid sequences located on the
XX human chromosome 13q31-q33 locus. The nucleotide sequences contain
XX biallelic markers and polymorphisms. Sequences AAH51602 - AAH51626 and
XX AAH62907 - AAH62915 represent cDNA human sbg1 cDNA sequences and protein
XX products. AAH51627 - AAH51631 and AAH62916 - AAH62918 represent g35018
XX cDNA sequences and protein products. Primers AAH51632 - AAH51699 are used
XX to isolate sbg1 cDNAs, while sbg1 exons from different primates are
XX represented by sequences AAH51642 - AAH51699. Nucleotide sequences of
XX amplicons which comprise biallelic markers located on the chromosome
XX 13q31-q33 locus are represented in AAH51700 - AAH51817. Biallelic markers
XX are represented in the sequences by degenerate/undefined base codes. PCR
XX primers AAH51818 and AAH51819 are used in the isolation of sequences of
XX the invention. The biallelic marker containing nucleotide sequences are
XX used to determine the identity of the nucleotide at a biallelic marker in
XX a sample DNA sequence. The nucleotide sequences may be labelled and used
XX for genotyping by determining the identity of a nucleotide at a Region D-
XX related biallelic marker in a biological sample from single or multiple
XX subjects. By determining the frequency of a biallelic marker in a
XX population an association between a genotype and a trait, a haplotype and
XX a trait and a phenotype and a trait can be detected. The sequences can be
XX used to determine a predisposition to or early onset of schizophrenia or
XX bipolar disorder or a beneficial response to or side effects related to
XX treatment against schizophrenia or bipolar disorder
XX
SQ Sequence 319608 BP; 101600 A; 56677 C; 58335 G; 102722 T; 0 U; 274 Other;
XX
XX Query Match 61.3%; Score 18.4; DB 3; Length 319608;
XX Best Local Similarity 78.6%; Pred. No. 3.8e+02;
XX Matches 22; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
XX
QY 1 CTGAGCCCTCTCGACTCGAGACTTCCG 28
DB 8690 CTGAGCCCTCTCGACTTGGAGATTACG 8717
XX
RESULT 12
AAS09301
XX AAS09301 standard; DNA; 319608 BP.
XX
AC AAS09301;
XX
XX 26-SEP-2001 (first entry)
XX
DE Human schizophrenia associated gene g35030 and biallelic markers A1-A71.
XX
XX Human; g35030; biallelic marker; A1-A71; chromosome 13q31-q33;
XX schizophrenia; bipolar disorder; ds.
XX
OS Homo sapiens.
XX
XX
XX Key Location/Qualifiers
XX FH 7938 .7958
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XX FT 8297 .8315
XX FT /*tag= b
XX FT /note= "binds primer 99-27943-150.mis"
XX FT 8304 .8328
XX FT /*tag= c
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XX FT 8316
XX FT /*tag= d
XX FT /note= "Biallelic marker A1"
XX FT

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FT	misc_binding	/note= "Binds primer 99-27935-193. mis"	FT		/note= "Binds primer 99-24639-163. mis"
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FT		/tag= k	FT	primer_bind	/tag= ai
FT	primer_bind	/note= "Binds primer 99-27935-193. mis complement"	FT		/note= "Binds primer 99-24639-163. mis complement"
FT		complement(21845. .21864)	FT	primer_bind	160770. .160787
FT		/tag= l	FT		/tag= aj
FT	primer_bind	/note= "Binds primer 99-27935. pu complement"	FT	primer_bind	/note= "Binds primer 99-24634. pu"
FT		65463. .65471	FT		complement(160785. .160802)
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FT	misc_binding	/note= "Binds primer 99-31960-363. mis"	FT		168962. .168986
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FT	primer_bind	/note= "Binds primer 99-24656. pu"	FT	primer_bind	170791. .170809
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FT		/tag= z	FT	misc_binding	/note= "Binds primer 99-16100-147. mis"
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FT primer_bind 173339..173357
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FT /note= "Binds primer 99-5862-167.mis"
FT misc_binding 173346..173370
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FT misc_feature 173358
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Query Match 61.3%; Score 18.4; DB 5; Length 319608;
* Best Local Similarity 78.6%; Pred. No. 3.8e+02;
Matches 22; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
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Qy 1 CTGACCCCTCTGACTCGAGTCCG 28
    ||||| ||||| ||||| |||||
Gb 8690 CTGACCCATCTCGATTGAGATTACG 8717
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RESULT 13
AAL52198
ID AAL52198 standard; cDNA; 340449 BP.
XX
XX AAL52198;
XX
XX 22-SEP-2003 (first entry)
XX
XX Human secreted protein genomic DNA coding sequence.
XX
XX Human; gene; ds; secreted protein; chromosome 5; tissue typing;
XX secreted protein-related disease; transgenic animal; drug screening;
XX pharmacogenomic analysts; single nucleotide polymorphism; SNP.
XX
XX Homo sapiens.
XX
XX Key
XX location/Qualifiers
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Best Local Similarity 87.0%; Pred. No. 4.7e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Oy      2 TGAACCCCTTCGACTCGAGACT 24
Db      25999 TGAACCCCTCCCACTCCAGAGT 26021

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RESULT 14
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ID      ADB74382 standard; DNA; 42325 BP.
XX      AC
XX      ADB74382;
XX      DT
XX      04-DEC-2003 (first entry)
XX      DE
XX      Mycobacterium leprae DNA #16.
XX      KW
XX      Non-naturally occurring peptide; anion pump protein; tuberculosis;
XX      hypersensitivity reaction; tuberculostatic; gene; ds.
XX      OS
XX      Mycobacterium leprae.
XX      PN
XX      US6583266-B1.
XX      PD
XX      24-JUN-2003.
XX      PF
XX      16-SEP-1994; 94US-00311731.
XX      PR
XX      19-AUG-1993; 93US-00109181.
XX      PR
XX      22-OCT-1993; 93US-00142558.
XX      PA
XX      (GENO-) GENOME THERAPEUTICS CORP.
XX      PI
XX      Smith DR, Mao J;
XX      DR
XX      WPI; 2003-656441/62.
XX      PS
XX      Disclosure; SEQ ID NO 131; 26pp; English.
XX      CC
XX      The invention relates to a non-naturally occurring peptide of
XX      Mycobacterium tuberculosis comprising an amino acid sequence
XX      corresponding to an anion pump protein. The invention also relates to a
XX      non-naturally occurring nucleic acid corresponding to a DNA sequence of
XX      Mycobacterium tuberculosis or Mycobacterium leprae. The new peptide is
XX      useful as a vaccine against Mycobacterium tuberculosis or Mycobacterium
XX      leprae or for screening for new tuberculosis drugs. Purified proteins

```

CC derived from the sequences of the invention may elicit a specific immune  
 CC response. The peptide may also be used to detect hypersensitivity  
 CC reactions of individuals exposed to Mycobacterium tuberculosis or  
 CC Mycobacterium leprae. The proteins and peptides may be affixed to solid  
 CC supports to detect antibodies typical of hypersensitivity reactions, from  
 CC a patient's sera. This sequence represents Mycobacterium leprae DNA of  
 CC the invention. Note: The sequence data for this patent did not form part  
 CC of the printed specification but was obtained in electronic format  
 CC directly from USPTO at [seqdata.uspto.gov/sequence.html](http://seqdata.uspto.gov/sequence.html).

XX  
 SQ Sequence 42325 BP; 9673 A; 13128 C; 11330 G; 8194 T; 0 U; 0 Other;

Query Match 60.0%; Score 18; DB 10; Length 42325;  
 Best Local Similarity 80.8%; Pred. No. 5e+02;

Matches 21; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 5 ACCCTCTCGACTCGAGATTCGGCT 30  
 ||||| ||||| ||||| ||||| |||||

DB 688 ACCCCACTCGACTCGACAGTACGCT 713

# RESULT 15

ADM27081\_10/c

Continuation (11 of 17) of ADM27081 from base 1000001 (Hyperthermophile Methanopyrus kan

WP Sequence split into 17 fragments LOCUS ADM27081 Accession Adm27081

WP Fragment Name

Begin End.

WP	ADM27081_00	1	110000
WP	ADM27081_01	100001	210000
WP	ADM27081_02	200001	310000
WP	ADM27081_03	300001	410000
WP	ADM27081_04	400001	510000
WP	ADM27081_05	500001	610000
WP	ADM27081_06	600001	710000
WP	ADM27081_07	700001	810000
WP	ADM27081_08	800001	910000
WP	ADM27081_09	900001	1010000
WP	ADM27081_10	1000001	1110000
WP	ADM27081_11	1100001	1210000
WP	ADM27081_12	1200001	1310000
WP	ADM27081_13	1300001	1410000
WP	ADM27081_14	1400001	1510000
WP	ADM27081_15	1500001	1610000
WP	ADM27081_16	1600001	1694968

Query Match 60.0%; Score 18; DB 11; Length 110000;

Best Local Similarity 80.8%; Pred. No. 5.4e+02;

Matches 21; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

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 ||||| ||||| ||||| ||||| |||||

DB 69684 CTTTGCCGCTCTCGACTCGAGAGTTC 69659

Search completed: April 26, 2005, 11:11:26  
 Job time : 433 secs